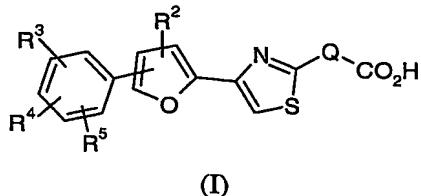


**CLAIMS:**

1. A compound of formula (I) or a pharmaceutically acceptable salt or prodrug thereof:

5



wherein

- 10      Q is  $(CH_2)_m[CH(R^1)]_n(CH_2)_p$ , where n is 0 or 1, and m and p are independently 0, 1 or 2;  
 R<sup>1</sup> is hydrogen, C<sub>1-6</sub> alkyl, C<sub>2-6</sub> alkenyl or C<sub>3-6</sub> alkynyl;
- 15      R<sup>2</sup> is hydrogen, halogen, C<sub>1-6</sub> alkyl optionally substituted by hydroxy or C<sub>1-6</sub> alkoxy, or phenyl  
 optionally substituted by one or more substituents selected from halogen, C<sub>1-6</sub> alkyl, CF<sub>3</sub>, OCF<sub>3</sub>, OR<sup>6</sup>, CN and methylenedioxo;
- 20      R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> are independently hydrogen, halogen, C<sub>1-6</sub> alkyl optionally substituted by hydroxy or C<sub>1-6</sub> alkoxy, CF<sub>3</sub>, OR<sup>6</sup>, COR<sup>7</sup>, NHCOR<sup>8</sup>, NHCONHR<sup>8</sup>, NHSO<sub>2</sub>R<sup>8</sup>, CONHR<sup>9</sup>, CN, SO<sub>2</sub>R<sup>8</sup> or NR<sup>10</sup>R<sup>11</sup>;  
 R<sup>6</sup> is hydrogen, C<sub>2-6</sub> alkenyl, C<sub>3-6</sub> alkynyl, C<sub>1-6</sub> alkyl optionally substituted by hydroxy or C<sub>1-6</sub> alkoxy, aryl or heteroaryl wherein aryl or heteroaryl is optionally substituted by one or more substituents selected from halogen, CF<sub>3</sub>, OCF<sub>3</sub>, CN, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy and methylenedioxo;
- 25      R<sup>7</sup> is C<sub>1-6</sub> alkyl, OR<sup>6</sup> or phenyl optionally substituted by one or more substituents selected from halogen, CF<sub>3</sub>, OCF<sub>3</sub>, CN, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy and NHCOR<sup>8</sup>;  
 R<sup>8</sup> is C<sub>1-6</sub> alkyl, C<sub>2-6</sub> alkenyl or C<sub>1-6</sub> alkoxy any of which may be optionally substituted by aryl or heteroaryl, wherein the aryl or heteroaryl is optionally substituted by one or more substituents selected from halogen, CF<sub>3</sub>, OCF<sub>3</sub>, OR<sup>6</sup>, CN, C<sub>1-6</sub> alkyl, methylenedioxo and NR<sup>10</sup>R<sup>11</sup>; C<sub>3-6</sub> cycloalkyl wherein the ring may contain up to two heteroatoms selected from NR<sup>12</sup>, S and O; or aryl or heteroaryl wherein the aryl or heteroaryl is optionally substituted by one or more substituents selected from halogen, CF<sub>3</sub>, OCF<sub>3</sub>, OR<sup>6</sup>, CN, C<sub>1-6</sub> alkyl, methylenedioxo and NR<sup>10</sup>R<sup>11</sup>;
- 30      R<sup>9</sup> is C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkylphenyl or phenyl, wherein alkyl may be interrupted by oxygen and wherein phenyl is optionally substituted by one or more substituents selected from halogen, C<sub>1-6</sub> alkyl, CF<sub>3</sub>, OCF<sub>3</sub>, CN, C<sub>1-6</sub> alkoxy and methylenedioxo;
- 35      R<sup>10</sup> and R<sup>11</sup> are independently hydrogen or C<sub>1-6</sub> alkyl, or together with the nitrogen to which they are attached form a 5- to 6-membered heterocyclic group which optionally contains an additional heteroatom selected from NR<sup>12</sup>, O and S; and  
 R<sup>12</sup> is hydrogen or C<sub>1-6</sub> alkyl;  
 provided that the compound is not:  
 i) 2-[4-[5-(2,4-dichlorophenyl)furan-2-yl]-1,3-thiazol-2-yl]acetic acid.
2. A compound according to claim 1 wherein Q is CH<sub>2</sub>.

3. A compound according to claim 1 or 2 wherein R<sup>2</sup> is hydrogen or halogen.
4. A compound according to any one of the preceding claims wherein R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> are independently, hydrogen, halogen, C<sub>1-6</sub> alkyl optionally substituted by hydroxyl or C<sub>1-6</sub> alkoxy, CF<sub>3</sub>, OR<sup>6</sup>, NHCOR<sup>8</sup> or CONHR<sup>9</sup>, wherein at least one of R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> is other than hydrogen.
5. A compound according to any one of the preceding claims wherein one of R<sup>3</sup> and R<sup>4</sup> is NHCOR<sup>8</sup> and the other is hydrogen or halogen and R<sup>5</sup> is hydrogen.
- 10 6. A compound according to any one of the preceding claims wherein R<sup>8</sup> is C<sub>1-6</sub> alkyl, C<sub>2-6</sub> alkenyl or C<sub>1-6</sub> alkoxy any of which may be optionally substituted by phenyl wherein the phenyl is optionally substituted by one or more substituents selected from halogen, CF<sub>3</sub>, OCF<sub>3</sub>, OR<sup>6</sup>, CN, C<sub>1-6</sub> alkyl, methylenedioxo and NR<sup>10</sup>R<sup>11</sup>; C<sub>3-6</sub> cycloalkyl wherein the ring may contain up to two heteroatoms selected from NR<sup>12</sup>, S and O; phenyl optionally substituted by one or more substituents selected from halogen, C<sub>1-6</sub> alkyl, CF<sub>3</sub>, OCF<sub>3</sub>, OR<sup>6</sup>, CN and methylenedioxo; or a 5- to 10-membered mono- or bicyclic heteroaryl group containing up to three heteroatoms selected from O, N and S which heteroaryl group may be substituted by C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy or halogen.
- 15 7. A compound according to claim 6 wherein R<sup>8</sup> is C<sub>1-6</sub> alkyl or C<sub>2-6</sub> alkenyl either of which may be optionally substituted by phenyl wherein the phenyl is optionally substituted by one or more substituents selected from halogen, CF<sub>3</sub>, OCF<sub>3</sub>, OR<sup>6</sup>, CN, C<sub>1-6</sub> alkyl, methylenedioxo and NR<sup>10</sup>R<sup>11</sup>; phenyl optionally substituted by one or more substituents selected from halogen, C<sub>1-6</sub> alkyl, CF<sub>3</sub>, OCF<sub>3</sub>, OR<sup>6</sup>, CN and methylenedioxo; or a 5- to 10-membered mono- or bicyclic heteroaryl group containing up to three heteroatoms selected from O, N and S which heteroaryl group may be substituted by C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy or halogen.
- 20 8. A compound of formula (I) as described in any one of Examples 1 to 24 or a pharmaceutically acceptable salt or prodrug thereof.
- 30 9. A compound selected from:  
2-[4-[5-[2-Chloro-4-[(2,4-dichlorophenylcarbonyl)amino]phenyl]furan-2-yl]-1,3-thiazol-2-yl]acetic acid,  
2-[4-[5-[2-Chloro-4-[3-(4-bromo)phenylacryloylamino]phenyl]furan-2-yl]-1,3-thiazol-2-yl]acetic acid,  
2-[4-[5-[2-Chloro-4-[3-(2,4-dichloro)phenylacryloylamino]phenyl]furan-2-yl]-1,3-thiazol-2-yl]acetic acid,  
2-[4-[5-[2-Chloro-4-[3-(3,5-difluoromethyl)phenylacryloylamino]phenyl]furan-2-yl]-1,3-thiazol-2-yl]acetic acid,  
2-[4-[5-[2-Chloro-4-(3-phenylacryloylamino)phenyl]furan-2-yl]-1,3-thiazol-2-yl]acetic acid,
- 35 40

2-[4-[5-[2-Chloro-4-[(4-trifluoromethoxyphenylcarbonyl)amino]phenyl]furan-2-yl]-1,3-thiazol-2-yl]acetic acid,

2-[4-[5-[2-Chloro-4-[(benzothiophene-2-carbonyl)amino]phenyl]furan-2-yl]-1,3-thiazol-2-yl]acetic acid,

5 2-[4-[5-[2-Chloro-4-[(6-chloro-4H-chromene-3-carbonyl)amino]phenyl]furan-2-yl]-1,3-thiazol-2-yl]acetic acid,

2-[4-[5-[2-Chloro-4-[(3,4-dichlorophenylcarbonyl)amino]phenyl]furan-2-yl]-1,3-thiazol-2-yl]acetic acid,

10 2-[4-[5-[2-Chloro-4-[(3-methoxyphenylcarbonyl)amino]phenyl]furan-2-yl]-1,3-thiazol-2-yl]acetic acid, and

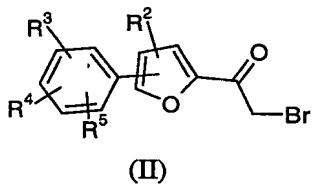
2-[4-[5-[2-Chloro-4-[(4-trifluoromethoxyphenylcarbonyl)amino]phenyl]furan-2-yl]-1,3-thiazol-2-yl]acetic acid,

and pharmaceutically acceptable salts and prodrugs thereof.

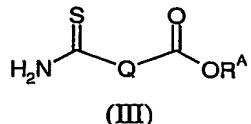
15 10. A compound according to any one of claims 1 to 9, without proviso i), for use in medicine.

11. A process for the preparation of a compound according to any one of claims 1 to 9 which comprises:

reacting a compound of formula (II):



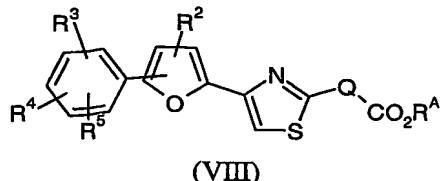
wherein R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> are as defined in claim 1, with a compound of formula (III):



25 wherein Q is as defined in claim 1 and R<sup>A</sup> is H, C<sub>1-6</sub> alkyl or a suitable protecting group; followed, where required, by deprotection of the group OR<sup>A</sup> to give the corresponding carboxylic acid.

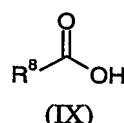
30 12. A process for the preparation of a compound according to any one of claims 1 to 9 wherein one or more of R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> is NHCOR<sup>8</sup> which comprises:

reacting a compound of formula (VIII):



wherein one or more of R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> is NH<sub>2</sub>, R<sup>2</sup> and Q are as defined in claim 1 and R<sup>A</sup> is as defined in claim 11, with a compound of formula (IX):

5



wherein R<sup>8</sup> is as defined in claim 1, in an amide bond formation reaction.

10 13. A pharmaceutical formulation comprising a compound according to any one of claims 1 to 9, without proviso i), together with a pharmaceutically acceptable carrier or excipient.

14. The use of a compound according to any one of claims 1 to 9, without proviso i), in the manufacture of a medicament for the inhibition of heparanase.

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15. The use of a compound according to any one of claims 1 to 9, without proviso i), in the manufacture of a medicament for the treatment of cancer.

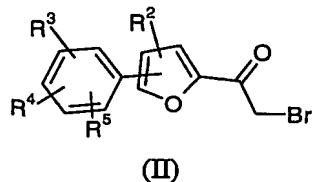
16. The use as claimed in claim 15 wherein the cancer is:

- 20 (a) a metastatic tumour cell type, such as, melanoma, lymphoma, leukaemia, fibrosarcoma, rhabdomyosarcoma, and mastocytoma; or  
 (b) a carcinoma, such as, colorectal cancer, prostate cancer, small cell lung cancer and non-small cell lung cancer, breast cancer, pancreatic cancer, bladder cancer, renal cancer, gastric cancer and ovarian cancer.

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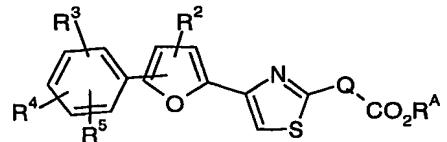
17. The use of a compound according to any one of claims 1 to 9, without proviso i), in the manufacture of a medicament for the treatment of a disease selected from angiogenesis or an angiogenesis dependent disease, an inflammatory disease, an autoimmune disease and a cardiovascular disease.

30 18. A compound of formula (II):



wherein, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> are as defined in claim 1.

19. A compound of formula (X):



(X)

5 wherein Q and R<sup>2</sup> are as defined in claim 1, R<sup>A</sup> is as defined in claim 11, at least one of R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> is NO<sub>2</sub> and the remainder are as defined in claim 1.